50 kDa —	Locus	UniquePep	Peptid	SpecCou	NSAF	Coverag	ProteinScore	Description
50 KDa —		tideCount	eCoun	nt		e		
0 kDa —			t					
50 KDa —	P12346	16	16	31	0.038478127	0.2865	65.6416	Serotransferrin
111.	P14480	14	14	27	0.048835529	0.1962	59.7884	Fibrinogen beta chain
0 kDa —	P01946	11	11	31	0.18913896	0.662	83.3401	Hemoglobin subunit alpha-1/2
75 kDa —	P26453	7	7	14	0.031261079	0.1495	27.2829	Basigin
	A0A0G2JSW3	7	16	36	0.212174317	0.7415	78.8489	Globin a4
50 kDa —	COJPT7	7	7	8	0.002626384	0.0428	21.4835	Filamin A
	P02680	6	6	11	0.021416097	0.1551	23.2189	Fibrinogen gamma chain
37 kDa —	Q7TQ70	6	6	15	0.016618513	0.0665	36.9458	Ac1873
	Q07936	6	6	8	0.020445509	0.2094	18.359	Annexin A2
	P07150	6	6	9	0.022535856	0.1532	18.8961	Annexin A1
kDa —	P09650	4	4	6	0.019993349	0.1769	13.9336	Mast cell protease 1
kDa —	G3V6P7	3	3	3	0.001326089	0.0158	5.1914	Myosin, heavy polypeptide 9, nor
kDa —	P10252	2	2	4	0.014439641	0.125	9.9145	CD48 antigen
	P31720	2	2	3	0.010608716	0.1061	6,2488	Complement C1q subcomponent
kDa <u>—</u>	101/10			5	0.010000710	0.1001	0.2400	subunitA
DOLLARS .	P20059	2	2	2	0.003766863	0.0413	4.0146	Hemopexin
	P00507	2	2	2	0.004029667	0.0395	4.9357	Aspartate aminotransferase, mitochondrial

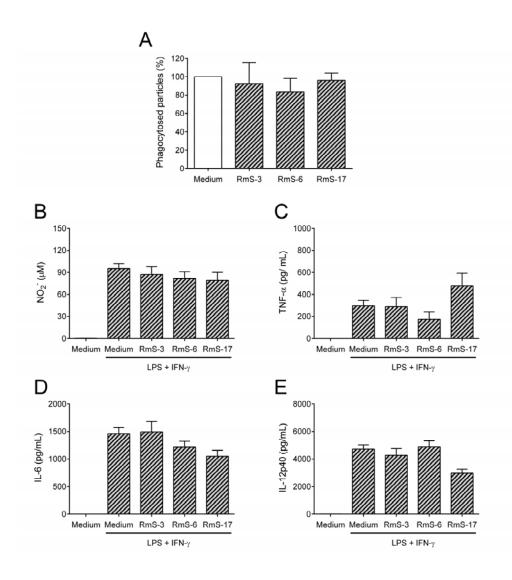
С

>P09650 Mast cell protease 1 MQALLFLMALLLPSGAGAEEIIGGVESRPH SRPYMAHLEITTERGYKATCGGFLVTRQF VMTAAHCKGRETTVTLGVHDVSKTESTQQKIKVEKQIVHPNYNFYSNLHDIMLLKLQKKA KVTPAVDVIPLPQPSDFLKPGKMCRAAGWGQTGVTKPTSNTLREVKQRIMDKEACKNYFH YNYNFQVCVGSPRKIRSAYKGDSGGPLVCAGVAHGIVSYGRGDAKPPAVFTRISPYVPWI NKVIKGKDLTSLSLHESESPS

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2 Supplementary Fig. S1. Native rat mast cell protease 1 (rMCP-1) is the main chymotrypsin-like 3 protease in peritoneum-derived cell extract. (A) Zymography was performed using 12% SDS-4 PAGE containing 0.1% casein. Gel was loaded with the protein extract containing chymotrypsin-5 like activity isolated from peritoneum-derived mast cells. (B) Protein extract containing 6 chymotrypsin-like activity isolated from peritoneum-derived mast cells was separated by 12% 7 SDS-PAGE following by Coomassie Blue staining. Band spot representing proteolytic activity 8 on zymography was excised and subjected to trypsin digestion followed by LC-MS/MS analysis. 9 (C) rMCP-1 sequence, highlighted peptides were identified by LC-MS/MS in the gel band with 10 proteolytic activity (4 unique to rMCP-1 with 17% coverage).

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13 Supplementary Fig. S2. Rhipicephalus microplus serpins do not affect phagocytosis or the 14 production of inflammatory mediators by murine macrophages. For the phagocytosis assay, cells 15 were incubated with medium only or in presence of each serpin (500 nM) followed by zymosan 16 particles opsonized with mouse serum. Phagocytosis was evaluated by light microscopy and the 17 percentage of phagocytosis was determined by the number of macrophages that had three or 18 more zymosan particles in each 100 cells (A). For the production of inflammatory mediators, 19 cells were preincubated with medium only or with each serpin (1000 nM each) for 1 h and 20 stimulated or not with LPS plus IFN-y (final concentration: 10 ng/ml of each). Cell culture 21 supernatants were collected NO determination by Griess reaction (B), and TNF- α (C), IL-6 (D) 22 and IL-12p40 (E) determination by ELISA.

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